

Josef Finsterer

List of Publications by Year in descending order

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Version: 2024-02-01

525
papers

6,118
citations

108046

37
h-index

111975

67
g-index

529
all docs

529
docs citations

529
times ranked

6244
citing authors

#	ARTICLE	IF	CITATIONS
1	Stroke-like episode of the optic nerve. Canadian Journal of Ophthalmology, 2024, 59, e77-e78.	0.4	5
2	Do seizures in POLG1-related mitochondrial disorder become refractory due to mitochondrion-toxic anti-seizure drugs?. Seizure: the Journal of the British Epilepsy Association, 2023, 104, 39-40.	0.9	0
3	Guillain-Barré syndrome is immunogenic in SARS-CoV-2 infected. Journal of Medical Virology, 2022, 94, 22-23.	2.5	3
4	Clinico-Genetic Spectrum of POLG1 Mutation Carriers from India. Journal of Molecular Neuroscience, 2022, 72, 45-47.	1.1	1
5	Lobar bleeding with ventricular rupture shortly after first dosage of an mRNA-based SARS-CoV-2 vaccine. Brain Hemorrhages, 2022, 3, 26-28.	0.4	5
6	Neurological side effects of SARS-CoV-2 vaccinations. Acta Neurologica Scandinavica, 2022, 145, 5-9.	1.0	106
7	Fatal status epilepticus: the broad phenotypic heterogeneity of NARS2 variants. Neurogenetics, 2022, 23, 73-74.	0.7	0
8	Real-world safety data for the Pfizer BNT162b2 SARS-CoV-2 vaccine, historical cohort study' by Shasha et al. Clinical Microbiology and Infection, 2022, 28, 452-453.	2.8	1
9	Comment on "Long-term results of palpebral fissure transfer with no lower eyelid spacer in chronic progressive external ophthalmoplegia". American Journal of Ophthalmology, 2022, 236, 319.	1.7	0
10	Broadening the diagnostic approach for SARS-CoV-2 associated myopathy and rhabdomyolysis. Therapeutic Apheresis and Dialysis, 2022, 26, 669-670.	0.4	0
11	Impaired hearing following SARS-CoV-2 vaccinations. International Journal of Infectious Diseases, 2022, 115, 215-216.	1.5	1
12	THE MICROBIOTA IN PARKINSON'S DISEASE: RANKING THE RISK OF HEART DISEASE. Annals of Nutrition and Metabolism, 2022, , .	1.0	0
13	The spectrum of neuroCOVID is broadening. Clinical and Experimental Neuroimmunology, 2022, 13, 127-128.	0.5	1
14	Anticoagulated de novo atrial flutter complicated by transitory ischemic attack in fatal COVID-19. Clinical Case Reports (discontinued), 2022, 10, e05246.	0.2	3
15	Affection of Cranial Nerves in COVID-19 Patients Should Prompt Suspicion of Guillain-Barre Syndrome. European Neurology, 2022, , 1-2.	0.6	0
16	Post-SARS-CoV-2 vaccination facial palsy requires extensive work-up and appropriate treatment. Indian Journal of Ophthalmology, 2022, 70, 346.	0.5	1
17	Consider differentials before diagnosing COVID-19 associated polyradiculitis. European Journal of Translational Myology, 2022, 32, .	0.8	6
18	Comments on "CSF-Confirmed SARS-CoV-2 Acute Encephalitis: SARS-CoV-2-Associated Encephalitis Is		

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19	The Broad Spectrum of Neuro-Radiological Abnormalities in Patients Infected with SARS-CoV-2 Supports the Diagnosis of Neuro-COVID-19. Korean Journal of Radiology, 2022, 23, 150.	1.5	2
20	Secondary mechanisms by which SARS-CoV-2 affects the brain. Revista Brasileira De Psiquiatria, 2022, , .	0.9	0
21	The etiology of SARS-CoV-2 associated intra-cranial hemorrhage is broad. Brain Hemorrhages, 2022, , .	0.4	0
22	MELAS with multiple stroke-like episodes due to the variant m.13513G>A in <i>MT-ND5</i>. Clinical Case Reports (discontinued), 2022, 10, e05361.	0.2	4
23	Sudden unexpected death in Parkinson's disease: Insights from clinical practice. Clinics, 2022, 77, 100001.	0.6	5
24	Nerve conduction studies support the classification of SARS-CoV-2 associated Guillain-Barre subtypes. Clinical and Experimental Neuroimmunology, 2022, 13, 95-96.	0.5	2
25	Exclude differentials before diagnosing SARS-CoV-2 associated acute, hemorrhagic, necrotising encephalitis. International Journal of Infectious Diseases, 2022, , .	1.5	0
26	Cognitive aspects of MELAS and CARASAL. Cerebral Circulation - Cognition and Behavior, 2022, 3, 100139.	0.4	0
27	Diagnose Aseptic Meningitis Caused by SARS-CoV-2 Vaccination Only After Ruling Out All Possible Differentials. Infection and Chemotherapy, 2022, 54, 185.	1.0	1
28	Ischemic stroke in 455 COVID-19 patients. Clinics, 2022, 77, 100012.	0.6	12
29	Acute Diffusion MRI Findings in Metabolic Encephalopathies are Diverse. Korean Journal of Radiology, 2022, 23, 381.	1.5	0
30	Unvaccinated patients may still experience SARS-CoV-2 associated polyradiculitis. Journal of Family Medicine and Primary Care, 2022, 11, 815.	0.3	0
31	Pathophysiological aspects of neuro-COVID. Revista Da Sociedade Brasileira De Medicina Tropical, 2022, 55, e0381.	0.4	0
32	Exacerbation of diabetic neuropathy can mimic SARS-CoV-2 associated pure dysautonomic Guillain-Barre syndrome. Journal of NeuroVirology, 2022, , .	1.0	0
33	Pattern Recognition in Mitochondrial Leukodystrophies is Hampered by the Peculiarities of Mitochondrial Genetics. American Journal of Neuroradiology, 2022, 43, E12-E12.	1.2	0
34	Neuro-COVID due to response against the virus. Neurologia I Neurochirurgia Polska, 2022, 56, 103-104.	0.6	0
35	Seizure and neuropsychiatric compromise as onset of SARS-CoV-2 associated encephalitis. Psychiatry Research Communications, 2022, 2, 100029.	0.2	2
36	Ischemic Stroke in a SARS-CoV-2-Positive Octagenarian Without Cardiovascular Risk Factors: A Case Report. Cureus, 2022, 14, e23654.	0.2	0

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37	Polyradiculitis and encephalomyelitis in the same patient following a SARS-CoV-2 vaccination. <i>Neurological Research and Practice</i> , 2022, 4, 11.	1.0	0
38	Consider alternative causes before allocating acute ischaemic stroke to COVID-19. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2022, , jnnp-2022-329029.	0.9	0
39	SARS-CoV-2â€“associated Guillainâ€“Barre syndrome requires extensive pre- and post-mortem examinations. <i>Journal of NeuroVirology</i> , 2022, , 1.	1.0	0
40	Metformin in m.3243A>G carriers can be both detrimental and beneficial. <i>Journal of Diabetes and Its Complications</i> , 2022, 36, 108184.	1.2	1
41	Bone quality in Duchenne muscular dystrophy. <i>Journal of Endocrinological Investigation</i> , 2022, , 1.	1.8	0
42	Facial palsy 12Âh after a first Moderna jab requires pathophysiological disclosure and verification of causality. <i>Clinical Imaging</i> , 2022, 83, 186-187.	0.8	1
43	Diagnosing SARS-CoV-2 vaccination associated rhombencephalitis requires comprehensive work-up and exclusion of differentials. <i>Neurological Research and Practice</i> , 2022, 4, 10.	1.0	2
44	Fatigue and Exercise Intolerance as Initial Manifestations of a Nonsyndromic Mitochondrial Disorder Due to the Variant m.3243A>G. <i>Case Reports in Neurological Medicine</i> , 2022, 2022, 1-3.	0.3	0
45	Neurological complications of COVID-19 in pediatric patients require comprehensive evaluation. <i>Journal of NeuroVirology</i> , 2022, , 1.	1.0	0
46	Real-world data about the side effects of SARS-CoV-2 vaccinations can be obtained only from representative samples undergoing comprehensive investigations. <i>Multiple Sclerosis and Related Disorders</i> , 2022, 60, 103696.	0.9	0
47	Diagnosing SARS-CoV-2 associated Guillain-Barre syndrome in children is challenging like in adults. <i>Annals of Medicine and Surgery</i> , 2022, 76, 103545.	0.5	0
48	Successful pregnancy in left ventricular hypertrabeculation/noncompaction with implanted cardioverter/defibrillator and a variant in the TPM1 Gen (c.425Aâ€“>â€“T) in mother and child. <i>Journal of Cardiology Cases</i> , 2022, , .	0.2	1
49	SARS-CoV-2 triggered relapse of multiple sclerosis. <i>Clinical Neurology and Neurosurgery</i> , 2022, 215, 107210.	0.6	11
50	Small fiber neuropathy underlying dysautonomia in COVIDâ€“19 and in postâ€“SARSâ€“CoVâ€“2 vaccination and longâ€“COVID syndromes. <i>Muscle and Nerve</i> , 2022, 65, .	1.0	7
51	When is the SARS-CoV-2 infection over and what is post-COVID?. <i>Annals of Medicine and Surgery</i> , 2022, 77, 103550.	0.5	0
52	Rule out differentials before blaming SARS-CoV-2. <i>Interdisciplinary Neurosurgery: Advanced Techniques and Case Management</i> , 2022, 29, 101551.	0.2	0
53	Do Patient/Doctor Benefits from Telephone/Electronic Contacts Exceed Those of Face-To-Face Visits?. <i>European Neurology</i> , 2022, 85, 77-78.	0.6	0
54	Phenotypic heterogeneity of Leigh syndrome due to <i>NDUFA12</i> variants is multicausal. <i>Human Mutation</i> , 2022, 43, 97-98.	1.1	1

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55	Venous sinus thrombosis after the second jab of an mRNA-based SARS-CoV-2 vaccine. <i>Brain Hemorrhages</i> , 2022, 3, 36-38.	0.4	9
56	Imaging abnormalities in pediatric neuro-COVID are more diverse than specified. <i>Biomedical Journal</i> , 2022, 45, 424-425.	1.4	4
57	SARS-CoV-2 vaccinations complicated by transverse myelitis. <i>Human Vaccines and Immunotherapeutics</i> , 2022, 18, 1-2.	1.4	3
58	Vaccine Adverse Event Reporting System Could Miss or Misinterpret Neurological Side Effects of COVID-19 Vaccinations. <i>Annals of Neurology</i> , 2022, 92, 157-158.	2.8	0
59	Guillain-Barré Syndrome Associated with COVID-19 Vaccination. <i>Emerging Infectious Diseases</i> , 2022, 28, 1079-1080.	2.0	1
60	Neuromyelitis optica complicating COVID vaccinations. <i>Multiple Sclerosis and Related Disorders</i> , 2022, 62, 103809.	0.9	2
61	Pathophysiology of SARS-CoV-2 associated ischemic stroke. <i>Journal of Medicine and Life</i> , 2022, 15, 149-150.	0.4	0
62	Immunodeficiency Should Be Excluded in Patients With Recurrent Viral Meningitis and Breakthrough COVID-19. <i>Journal of Korean Medical Science</i> , 2022, 37, e161.	1.1	0
63	Familial Intracranial Aneurysm Requires Not Only Whole-Exome Sequencing, But Also Mitochondrial DNA Sequencing. <i>Korean Journal of Radiology</i> , 2022, 23, 566.	1.5	1
64	Consider Transverse Myelitis as a Complication of a SARS-CoV-2 Vaccination. <i>Journal of Korean Medical Science</i> , 2022, 37, e150.	1.1	2
65	Neurological disease in pregnant females with COVID-19 may not only be attributable to SARS-CoV-2. <i>Acta Neurologica Scandinavica</i> , 2022, , .	1.0	2
66	De novo altered mental state after SARS-CoV-2 vaccination requires extensive diagnostic work-up. <i>Annals of Medicine and Surgery</i> , 2022, 77, 103724.	0.5	0
67	The diagnosis of SARS-CoV-2 associated ADEM requires the exclusion of all differential diagnoses. <i>Annals of Medicine and Surgery</i> , 2022, 77, 103662.	0.5	0
68	Intensification of an Essential Tremor by SARS-CoV-2. <i>Cerebellum</i> , 2022, , 1.	1.4	0
69	No need for brain biopsy in acute disseminated encephalomyelitis after first Sputnik-V jab. <i>Brain, Behavior, & Immunity - Health</i> , 2022, 22, 100464.	1.3	0
70	Wernicke Encephalopathy Mimicking MELAS. <i>Medicina (Lithuania)</i> , 2022, 58, 660.	0.8	1
71	Parkinson's Disease, Premature Mortality, and Amygdala. <i>Movement Disorders</i> , 2022, 37, 1110-1111.	2.2	1
72	Letter to the Editor: Propacetamol-Induced Rhabdomyolysis or COVID-Vaccine-Related Inflammatory Myopathy?. <i>Journal of Korean Medical Science</i> , 2022, 37, .	1.1	0

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73	The Spectrum of Renal Abnormalities in Mitochondrial Disorders Is Broad. <i>Kidney International Reports</i> , 2022, , .	0.4	1
74	Consider cerebral tuberculosis as differential of SARS-CoV-2-associated acute, haemorrhagic, necrotising encephalitis. <i>Egyptian Journal of Neurology, Psychiatry and Neurosurgery</i> , 2022, 58, .	0.4	0
75	Consider alternative causes of thrombo-embolism to SARS-CoV-2 in COVID-19 patients. <i>Annals of Medicine and Surgery</i> , 2022, 78, 103875.	0.5	0
76	Real world data speak a different language about the outcome of pregnancies undergoing SARS-CoV-2 vaccinations. <i>Annals of Medicine and Surgery</i> , 2022, 78, .	0.5	0
77	SARS-CoV-2 vaccinations reduce the prevalence of post-COVID Guillain-Barre syndrome. <i>Clinics</i> , 2022, 77, 100064.	0.6	3
78	Neuro-COVID is not at variance between children and adults. <i>European Journal of Paediatric Neurology</i> , 2022, , .	0.7	0
79	Side effects of SARS-CoV-2 vaccines should be assessed by unbiased professionals on-site. <i>Human Vaccines and Immunotherapeutics</i> , 2022, 18, .	1.4	1
80	Assessing the anesthetic effectiveness of remimazolam in MELAS patients requires careful investigations. <i>JA Clinical Reports</i> , 2022, 8, .	0.2	1
81	Diagnosing Weber syndrome requires compliance with diagnostic criteria and compatibility with cerebral imaging. <i>Annals of Medicine and Surgery</i> , 2022, , 104044.	0.5	0
82	Ischemic and Metabolic Stroke Can Co-occur in m.3243A>G Carriers: A Case Report. <i>Cureus</i> , 2022, , .	0.2	3
83	Determining prediction factors of post-neurosurgical thrombosis requires consideration of the entire spectrum of risk factors. <i>Annals of Medicine and Surgery</i> , 2022, 79, .	0.5	0
84	Do not miss Bickerstaff encephalitis as a complication of SARS-CoV-2 vaccines. <i>Radiology Case Reports</i> , 2022, 17, 2824-2825.	0.2	1
85	Letter to the Editor: Pre-Existing Neuropathy Favours SARS-CoV-2 Vaccination Associated Guillain-Barre Syndrome. <i>Journal of Korean Medical Science</i> , 2022, 37, .	1.1	1
86	Letter to the Editor: Finger Extensor Weakness Is Not a Novel Clinical Feature of SARS-CoV-2 Associated Guillain-Barre Syndrome. <i>Journal of Korean Medical Science</i> , 2022, 37, .	1.1	0
87	Comment on "MOG antibody-associated encephalomyelitis mimicking bacterial meningomyelitis following ChAdOx1 nCoV-19 vaccination: a case report"™. <i>Therapeutic Advances in Neurological Disorders</i> , 2022, 15, 175628642211063.	1.5	0
88	Retinal artery/vein occlusion complicating SARS-CoV-2 vaccinations. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2022, 31, 106617.	0.7	2
89	Carriers of POLG1 variants require investigations for multisystem disease and for mtDNA variations. <i>Neurological Research and Practice</i> , 2022, 4, .	1.0	0
90	Is Guillain Barre syndrome truly caused by SARS-CoV-2?. <i>American Journal of Emergency Medicine</i> , 2021, 45, 649.	0.7	1

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91	COVID-19 polyradiculitis in 24 patients without SARS-CoV-2 in the cerebrospinal fluid. <i>Journal of Medical Virology</i> , 2021, 93, 66-68.	2.5	25
92	Infectious/parainfectious, nonvascular, nonhypoxic central nervous system disease in 48 COVID-19 patients. <i>Journal of Medical Virology</i> , 2021, 93, 626-630.	2.5	0
93	Volume loss and altered neuronal composition in the brainstem reticular zone may not cause sudden unexpected death in epilepsy. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 171-172.	1.8	1
94	Kidney transplantation in m.3243A>G carriers has outcome implications. <i>CKJ: Clinical Kidney Journal</i> , 2021, 14, 723-724.	1.4	1
95	SARS-CoV-2-associated Guillain-Barre syndrome in 62 patients. <i>European Journal of Neurology</i> , 2021, 28, e10-e12.	1.7	33
96	SARS-CoV-2 myopathy. <i>Journal of Medical Virology</i> , 2021, 93, 1852-1853.	2.5	7
97	Letter to the editor: sudden death in Parkinson's disease: treating hypertension in the elderly is essential. <i>Expert Opinion on Pharmacotherapy</i> , 2021, 22, 1633-1634.	0.9	0
98	What the neuroradiologist should additionally consider in SARS-CoV-2 infection. <i>Emergency Radiology</i> , 2021, 28, 437-438.	1.0	0
99	In Reference to Impact of Fiberoptic Endoscopic Evaluation of Swallowing Outcomes and Dysphagia Management in Neurodegenerative Diseases. <i>Laryngoscope</i> , 2021, 131, E338.	1.1	0
100	Attributing increased prevalence of facial palsy to SARS-CoV-2 requires evidence. <i>Brain and Behavior</i> , 2021, 11, e01996.	1.0	5
101	Does SARS-CoV-2 truly cause infectious myopathy?. <i>Journal of the Formosan Medical Association</i> , 2021, 120, 1032-1033.	0.8	0
102	Myotoxic drugs and immunodeficiency may contribute to the poor outcome of COVID-19 patients with myotonic dystrophy. <i>Acta Neurologica Belgica</i> , 2021, 121, 799-800.	0.5	2
103	SARS-CoV-2 triggered Takotsubo in 38 patients. <i>Journal of Medical Virology</i> , 2021, 93, 1236-1238.	2.5	18
104	Energy requirements in m.3243A>G carriers depend on multiple factors. <i>Journal of Parenteral and Enteral Nutrition</i> , 2021, 45, 227-228.	1.3	0
105	Exclude hereditary and acquired differential disorders before attributing retinoschisis to Kears-Sayre syndrome. <i>Ophthalmic Genetics</i> , 2021, 42, 99-99.	0.5	0
106	Cardiac disease in mitochondrial disorders. <i>Heart Failure Reviews</i> , 2021, 26, 727-728.	1.7	0
107	Autonomic dysfunction may not be the only cause of SUDEP. <i>Acta Neurologica Scandinavica</i> , 2021, 143, 217-218.	1.0	0
108	Ischemic Stroke in COVID-19 Patients May Be Incidentally but Not Causally Related to the Infection. <i>Cerebrovascular Diseases</i> , 2021, 50, 361-362.	0.8	4

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109	Guillain-Barre syndrome 15 days after COVID-19 despite SARS-CoV-2 vaccination. IDCases, 2021, 25, e01226.	0.4	9
110	Prevention of Parkinson's disease-related sudden death. Clinics, 2021, 76, e3266.	0.6	2
111	Fatal SARS-CoV-2 Associated Rhabdomyolysis Requires Elucidation. Journal of Primary Care and Community Health, 2021, 12, 215013272110052.	1.0	0
112	Connectivity on fMRI in the MELAS brain may strongly depend on heteroplasmy and extension or dynamics of stroke-like lesions. NeuroImage: Clinical, 2021, 30, 102591.	1.4	0
113	Metabolic or ischemic stroke in succinic semi-aldehyde dehydrogenase deficiency due to the homozygous variant c. 1343 + 1_1343 + 3delGTAinsTT in ALDH5A1. Annals of Indian Academy of Neurology, 2021, 24, 303.	0.2	1
114	Parkinson-related neuropathy. Clinics, 2021, 76, e2675.	0.6	2
115	Sudden death in a patient with epilepsy and arterial hypertension: time for re-assessment. Clinics, 2021, 76, e3023.	0.6	1
116	Multifocal T2-/DWI-hyperintense cerebral lesions in COVID-19 not necessarily imply demyelination. Arquivos De Neuro-Psiquiatria, 2021, 79, 92-93.	0.3	1
117	Neuromuscular involvement in COVID-19 patients. Annals of Indian Academy of Neurology, 2021, 24, 768.	0.2	0
118	Mental compromise in SARS-CoV-2 infected patients is multicausal, organic or inorganic. Brain Communications, 2021, 3, fcab218.	1.5	0
119	Stroke-like lesions in mitochondrial disease may resemble ischemic stroke. Journal of Family Medicine and Primary Care, 2021, 10, 3151.	0.3	2
120	Cardiac disease in mitochondrial membrane protein-associated neurodegeneration (MPAN) due to variants in C19orf12. Parkinsonism and Related Disorders, 2021, 83, 13-14.	1.1	1
121	Perampanel may be beneficial in Leigh syndrome by its anti-oxidative but not anti-epileptic effect. Brain and Development, 2021, 43, 360.	0.6	1
122	More likely than through head trauma: is LHON triggered by mitochondrion-toxic drugs or oxidative stress. Documenta Ophthalmologica, 2021, 142, 395-396.	1.0	1
123	Takotsubo syndrome in COVID-19 requires elucidation of the pathophysiological background. Cardiovascular Revascularization Medicine, 2021, 34, 148-148.	0.3	0
124	Comment on "Mitochondrial Neurogastrointestinal Encephalomyopathy: Novel Pathogenic Mutation in Thymidine Phosphorylase Gene in a Patient from Cape Verde Islands". Case Reports in Neurological Medicine, 2021, 2021, 1-2.	0.3	0
125	Sequential Stroke-Like Lesions in MELAS are Common and Diagnosable upon Multimodal MRI. Canadian Journal of Neurological Sciences, 2021, 48, 744-744.	0.3	2
126	Neurologic manifestations of COVID-19. Polish Archives of Internal Medicine, 2021, 131, 208-208.	0.3	0

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127	Perspectives of Neuro-COVID: Myasthenia. <i>Frontiers in Neurology</i> , 2021, 12, 635747.	1.1	7
128	Is unilateral facial palsy truly caused by SARS-CoV-2?. <i>Arquivos De Neuro-Psiquiatria</i> , 2021, 79, 183-183.	0.3	1
129	Re. "To bee or not to bee? The bee extract propolis as a bioactive compound in the burden of lifestyle diseases" <i>Nutrition</i> , 2021, 93, 111241.	1.1	0
130	m.3243A>G carriers develop syndromic or non-syndromic multisystem phenotypes over time. <i>CEN Case Reports</i> , 2021, 10, 614-615.	0.5	2
131	SARS-CoV-2 associated polyradiculitis and myocarditis may favour Takotsubo syndrome. <i>Medical Hypotheses</i> , 2021, 148, 110509.	0.8	0
132	Brain and nerves affected before the lungs in COVID-19. <i>Acta Neurologica Scandinavica</i> , 2021, 143, 675-676.	1.0	3
133	The MELAS phenotype may not only be determined by heteroplasmy of causative mtDNA variants. <i>European Journal of Endocrinology</i> , 2021, 184, L5-L6.	1.9	3
134	Vascular Damage May Mimic Retinitis and Optic Neuritis in COVID-19. <i>Current Eye Research</i> , 2021, 46, 1934-1935.	0.7	11
135	MELAS can be delineated from CADASIL by genotype and phenotype. <i>Neurobiology of Aging</i> , 2021, 103, 128-129.	1.5	0
136	Involvement of the cardiac conduction system in Kearns-Sayre syndrome is progressive. <i>Europace</i> , 2021, 23, 979-980.	0.7	2
137	Superficial siderosis due to multiple cavernomas: an uncommon cause of early-onset dementia. <i>Psychogeriatrics</i> , 2021, 21, 434-437.	0.6	2
138	Prevent Hyperglycemia in MELAS by Measuring HbA1c Values and by Avoiding Triggering Events. <i>Pediatric Neurology</i> , 2021, 116, 60.	1.0	0
139	MuSK-positive myasthenia may be triggered not only by SARS-CoV-2. <i>European Journal of Neurology</i> , 2021, 28, e80-e81.	1.7	1
140	Coronavirus Disease 2019 Can Be Complicated by Immune-encephalopathy Rather Than Encephalitis. <i>Clinical Infectious Diseases</i> , 2021, 73, 1744-1744.	2.9	0
141	Fatalities of COVID-19 are rather attributable to multisystem inflammatory syndrome than infectious meningitis or sepsis. <i>Indian Journal of Medical Microbiology</i> , 2021, 39, 393-394.	0.3	0
142	Clinical and Pathophysiologic Spectrum of Neuro-COVID. <i>Molecular Neurobiology</i> , 2021, 58, 3787-3791.	1.9	25
143	Triggers of Takotsubo syndrome should be identified by exclusion. <i>Journal of Cardiac Surgery</i> , 2021, 36, 2184-2184.	0.3	0
144	Guillain-Barre syndrome in 220 patients with COVID-19. <i>Egyptian Journal of Neurology, Psychiatry and Neurosurgery</i> , 2021, 57, 55.	0.4	90

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145	Consider differentials before diagnosing SARS-CoV-2 associated Guillain-Barré syndrome. <i>Journal of Medical Virology</i> , 2021, 93, 5246-5247.	2.5	1
146	SARS-CoV-2 vaccines are not free of neurological side effects. <i>Acta Neurologica Scandinavica</i> , 2021, 144, 109-110.	1.0	26
147	Exacerbating Guillain-Barré Syndrome Eight Days after Vector-Based COVID-19 Vaccination. <i>Case Reports in Infectious Diseases</i> , 2021, 2021, 1-3.	0.2	36
148	Stroke-Like Lesion in an m.3243A>G Carrier Presenting as Hyperperfusion and Hypometabolism. <i>Cureus</i> , 2021, 13, e15487.	0.2	3
149	tRNA variants causing Leber's hereditary optic neuropathy?. <i>Irish Journal of Medical Science</i> , 2021, , 1.	0.8	0
150	SARS-CoV-2 associated Miller-Fisher syndrome or Polyneuritis cranialis. <i>Environmental Science and Pollution Research</i> , 2021, 28, 46082-46084.	2.7	1
151	Diagnose SARS-CoV-2 associated Guillain-Barré syndrome upon appropriate criteria and after exclusion of differentials. <i>Journal of Medical Virology</i> , 2021, 93, 5687-5688.	2.5	1
152	Workup of MERRF and MELAS fatalities requires autopsy of the brain and revision of intra-vitam investigations. <i>Pathology International</i> , 2021, 71, 559-560.	0.6	0
153	Spinal cord involvement in LHON requires pathophysiological clarification. <i>Multiple Sclerosis and Related Disorders</i> , 2021, 51, 102920.	0.9	0
154	SARS-CoV-2 associated rhabdomyolysis in 32 patients. <i>Turkish Journal of Medical Sciences</i> , 2021, 51, 1597-1600.	0.4	13
155	Is SARS-CoV-2 responsible for relapses of Parkinson's disease?. <i>Egyptian Journal of Neurology, Psychiatry and Neurosurgery</i> , 2021, 57, 90.	0.4	2
156	Neuropsychiatric disorders and COVID-19. <i>Lancet Psychiatry</i> , 2021, 8, 563-564.	3.7	0
157	Peripheral neuropathy in COVID-19 is due to immune-mechanisms, pre-existing risk factors, anti-viral drugs, or bedding in the Intensive Care Unit. <i>Arquivos De Neuro-Psiquiatria</i> , 2021, 79, 924-928.	0.3	46
158	Noncompaction and the novel variant c.425A>T in TPM1. <i>Acta Cardiologica</i> , 2021, , 1-2.	0.3	1
159	m.3243A>G Maculopathy. <i>Klinische Monatsblätter Fur Augenheilkunde</i> , 2021, 238, 827-827.	0.3	0
160	Do Not Replace Personal Patient/Doctor Interactions by Electronic Visits in Myotonic Dystrophy Patients. <i>European Neurology</i> , 2021, , 1-2.	0.6	2
161	Intracerebral bleeding after Janus-kinase inhibitor baricitinib for COVID-19. <i>Brain Hemorrhages</i> , 2021, 2, 151-152.	0.4	3
162	Infectious and immune-mediated central nervous system disease in 48 COVID-19 patients. <i>Journal of Clinical Neuroscience</i> , 2021, 90, 140-143.	0.8	14

#	ARTICLE	IF	CITATIONS
163	SARS-CoV-2 vaccinations are unsafe for those experiencing post-vaccination Guillain-Barre syndrome. <i>Annals of Medicine and Surgery</i> , 2021, 68, 102584.	0.5	5
164	Knowledge about the characteristics of stroke-like lesions is expandable. <i>Metabolic Brain Disease</i> , 2021, 36, 1697-1698.	1.4	0
165	COVID-19 associated cranial nerve neuropathy: A systematic review. <i>Bosnian Journal of Basic Medical Sciences</i> , 2021, , .	0.6	35
166	<sc>Cerebroâ€Spinalâ€Fluid</sc> Cytokine Profiles Do Not Reliably Delineate Encephalopathy and Inflammation in <sc>Neuroâ€COVID</sc>. <i>Annals of Neurology</i> , 2021, 90, 695-695.	2.8	0
167	Presentation and pathophysiology of neuro-COVID. <i>Drugs in Context</i> , 2021, 10, 1-2.	1.0	0
168	Neuro-COVID Requires Comprehensive Work-up. <i>Indian Journal of Critical Care Medicine</i> , 2021, 25, 956-957.	0.3	0
169	Survival from liver transplantation in mitochondrial disorders not only depends on periprocedural complications but also on the genotype and on mitochondrionâ€toxic drugs. <i>Pediatric Transplantation</i> , 2021, 25, e14137.	0.5	0
170	SARS-CoV-2 associated Guillain-Barre syndrome after awaking on the ICU: consider differentials. <i>Tuberkuloz Ve Toraks</i> , 2021, 69, 429-430.	0.2	0
171	Guillain-Barre syndrome is a definite complication of SARS-CoV-2. <i>Annals of Medicine and Surgery</i> , 2021, , 102800.	0.5	0
172	Symptomatic peduncular, cavernous bleeding following SARS-CoV-2 vaccination induced immune thrombocytopenia. <i>Brain Hemorrhages</i> , 2021, 2, 169-171.	0.4	12
173	SARS-CoV-2 in cardiomyocytes. <i>International Journal of Infectious Diseases</i> , 2021, 112, 195.	1.5	0
174	Pathophysiology of SARS-CoV-2-associated ischemic stroke is variegated. <i>Egyptian Journal of Neurology, Psychiatry and Neurosurgery</i> , 2021, 57, 120.	0.4	0
175	Reagibility of intracerebral arteries does not reflect dynamics of a stroke-like lesions. <i>Journal of Neuroradiology</i> , 2021, 48, 367-368.	0.6	0
176	Post-COVID Guillain-Barre syndrome: comparison of two reviews. <i>Canadian Journal of Neurological Sciences</i> , 2021, , 1-4.	0.3	0
177	Exclude differentials before attributing post-COVID fatigue to myopathy. <i>Clinical Neurophysiology</i> , 2021, 132, 2324-2325.	0.7	2
178	Diagnosing Myoclonic Epilepsy With Ragged-Red Fibers Syndrome Requires Documentation of a Causative Mutation. <i>A&A Practice</i> , 2021, 15, e01525.	0.2	0
179	Sinus venous thrombosis as a complication of COVID-19-associated hypercoagulability. <i>Egyptian Journal of Neurology, Psychiatry and Neurosurgery</i> , 2021, 57, 132.	0.4	1
180	Cortical cystic lesions â€“ A typical endpoint of a stroke-like lesion. <i>Brain and Development</i> , 2021, 43, 972-973.	0.6	1

#	ARTICLE	IF	CITATIONS
181	Suspect MELAS early to prevent unnecessary costs and burdens for the affected. Radiology Case Reports, 2021, 16, 3451-3452.	0.2	0
182	Sudden death in a rat model of Parkinson's disease. Clinics, 2021, 76, e2974.	0.6	0
183	Extrapulmonary onset manifestations of COVID-19. Clinics, 2021, 76, e2900.	0.6	29
184	Broken Heart (Takotsubo) Syndrome Triggered by SARS-CoV-2. SN Comprehensive Clinical Medicine, 2021, 3, 60-61.	0.3	0
185	Endometriosis as Initial Manifestation of Myotonic Dystrophy Type-2. Journal of Neurosciences in Rural Practice, 2021, 12, 219-221.	0.3	0
186	Antiviral activity of Brazilian Green Propolis extract against SARS-CoV-2 (Severe Acute Respiratory) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50	0.6	26
187	SARS-CoV-2 and myasthenia. Journal of Medical Virology, 2021, 93, 4133-4135.	2.5	3
188	Comment on peripheral polyneuropathy associated with COVID-19 in two patients: A musculoskeletal ultrasound case report. Journal of Medical Ultrasound, 2021, 29, 134.	0.2	0
189	COVID-19 polyradiculitis in 24 patients without SARS-CoV-2 in the cerebro-spinal fluid. , 2021, 93, 66.		2
190	Post SARS-CoV-2 vaccination Guillain-Barre syndrome in 19 patients. Clinics, 2021, 76, e3286.	0.6	41
191	Letter to the Editor: Ischemic Stroke of the Corpus Callosum after SARS-CoV-2 Vaccination. Journal of Korean Medical Science, 2021, 36, e288.	1.1	1
192	Impact of SARS-CoV-2 vaccines on the nervous system. Clinical Imaging, 2021, 82, 13-14.	0.8	1
193	First Reported Case of Reversible Cerebral Vasoconstriction Syndrome After a SARS-CoV-2 Vaccine. Cureus, 2021, 13, e19987.	0.2	11
194	Before attributing encephalomyelitis to SARS-CoV-2 vaccinations thoroughly exclude differentials. Annals of Clinical and Translational Neurology, 2021, 8, 2222-2223.	1.7	1
195	Rheumatoid arthritis: Propolis consumption can be useful. Journal of Food Biochemistry, 2021, 45, e14009.	1.2	0
196	Apparent onset of COVID-19 after onset of SARS-CoV-2 associated Guillain-Barre syndrome. Travel Medicine and Infectious Disease, 2021, 44, 102201.	1.5	0
197	Takotsubo rather than Kounis syndrome complicating SARS-CoV-2 vaccination. Journal of Emergency Medicine, 2021, , .	0.3	1
198	SARS-CoV-2-associated Guillain-Barre syndrome is not infrequent. Revista Da Associação Médica Brasileira, 2021, 67, 1521-1522.	0.3	0

#	ARTICLE	IF	CITATIONS
199	Post-SARS-CoV-2 vaccination venous sinus thrombosis: a literature review of 308 cases. Egyptian Journal of Neurology, Psychiatry and Neurosurgery, 2021, 57, 179.	0.4	3
200	Reply to the letter "Venous sinus thrombosis after the second dose of SARS-CoV-2 vaccine administration" by Mungmunpantip and Wiwanitkit. Brain Hemorrhages, 2021, , .	0.4	0
201	Discrepancy between Mild Muscle Pathology and Severe Muscular Compromise in COVID-19 Suggests Nonviral Etiologies. Journal of Neuropathology and Experimental Neurology, 2021, 80, 1088-1090.	0.9	0
202	The heart in m.3243A>G carriers. Herz, 2020, 45, 356-361.	0.4	18
203	NEMMLAS Due to Biallelic WARS2 Variants. Journal of Child Neurology, 2020, 35, 175-175.	0.7	0
204	Cerebellar stroke-like lesions in Leigh syndrome due to the variant m.8993T>C in MT-ATP6. ENeurologicalSci, 2020, 18, 100203.	0.5	0
205	Takotsubo syndrome in Duchenne muscular dystrophy may be triggered by epilepsy. Journal of Cardiology Cases, 2020, 21, 82.	0.2	1
206	Muscle biopsy is not diagnostic for MELAS. Journal of the Neurological Sciences, 2020, 410, 116670.	0.3	2
207	Cerebrospinal-fluid ATP is inappropriate as a biomarker of disease severity and treatment response in MELAS. Mitochondrion, 2020, 51, 140-141.	1.6	0
208	Low heteroplasmy rates in clinically affected m.3243A>G carriers not necessarily explain the phenotype. Journal of the Neurological Sciences, 2020, 409, 116614.	0.3	0
209	Comment on: Factors Affecting Generalization of Ocular Myasthenia Gravis in Patients With Positive Acetylcholine Receptor Antibodies. American Journal of Ophthalmology, 2020, 210, 193-194.	1.7	0
210	Viability of diffusion tensor imaging for assessing retro-chiasmatic involvement in Kearns-Sayre syndrome remains elusive. Neuroradiology, 2020, 62, 131-132.	1.1	0
211	Pathogenicity of the variant m.13376A>C in ND5 remains unproven. Journal of the Neurological Sciences, 2020, 409, 116615.	0.3	0
212	Stroke-like episodes in OPA1 carriers require comprehensive work-up and therapeutic considerations. Metabolic Brain Disease, 2020, 35, 253-254.	1.4	0
213	The Spectrum of Neuromuscular Disorders Admitted to a Pediatric Intensive Care Unit Is Broader Than Anticipated. Journal of Child Neurology, 2020, 35, 300-301.	0.7	1
214	Photosensitive Epilepsy and Polycystic Ovary Syndrome as Manifestations of MERRF. Case Reports in Neurological Medicine, 2020, 2020, 1-4.	0.3	5
215	Letter by Finsterer Regarding Article, "COVID-19-Associated Stress (Takotsubo) Cardiomyopathy": Circulation: Cardiovascular Imaging, 2020, 13, e011577.	1.3	1
216	SARS-CoV-2-associated Takotsubo is not necessarily triggered by the infection. IJC Heart and Vasculature, 2020, 30, 100606.	0.6	2

#	ARTICLE	IF	CITATIONS
217	Are stroke-like lesions in Leber's hereditary optic neuropathy mistaken as MS-like cerebral lesions?. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 45, 102321.	0.9	1
218	Propolis and coronavirus disease 2019 (COVID-19): Lessons from nature. <i>Complementary Therapies in Clinical Practice</i> , 2020, 41, 101227.	0.7	24
219	Is the variant m.9176T>C in <i>MT-ATP6</i> truly responsibly for Leigh syndrome?. <i>Pediatrics International</i> , 2020, 62, 1217-1217.	0.2	1
220	Outcome of endoscopic lipoma resection in myoclonic epilepsy with ragged-red fibers syndrome may depend on the genotype. <i>Asian Journal of Surgery</i> , 2020, 43, 1024-1025.	0.2	0
221	Cerebellar stroke-like lesions in Leigh syndrome may mimic cerebellar cortical bleeding. <i>ENeurologicalSci</i> , 2020, 19, 100234.	0.5	0
222	Secondary manifestations of mitochondrial disorders. <i>Journal of Zhejiang University: Science B</i> , 2020, 21, 590-592.	1.3	3
223	Bulbospinal muscular atrophy (Kennedy disease) responsive to immunoglobulins?. <i>Clinical Case Reports (discontinued)</i> , 2020, 8, 1223-1225.	0.2	1
224	Left Ventricular Noncompaction Syndrome: Genetic Insights and Therapeutic Perspectives. <i>Current Cardiology Reports</i> , 2020, 22, 84.	1.3	17
225	Optimising therapeutic strategies for acute stroke-like lesions in MELAS. <i>ENeurologicalSci</i> , 2020, 21, 100278.	0.5	2
226	SARS-CoV-2-associated critical ill myopathy or pure toxic myopathy?. <i>International Journal of Infectious Diseases</i> , 2020, 101, 56.	1.5	5
227	Factors Influencing Central Nervous System Abnormalities in m.11778G>A Carriers. <i>Brain Sciences</i> , 2020, 10, 513.	1.1	1
228	Consider Stroke-Like Episodes as a Differential of Migrainous Stroke. <i>Journal of Neurosciences in Rural Practice</i> , 2020, 11, 219-220.	0.3	0
229	Diagnosing myasthenic crisis in SARS-CoV-2 infected patients requires adherence to appropriate criteria. <i>Journal of the Neurological Sciences</i> , 2020, 417, 117062.	0.3	5
230	Prospective studies on the efficacy of rituximab for myasthenia gravis are warranted. <i>European Journal of Neurology</i> , 2020, 27, e95.	1.7	0
231	Multisystem disease including stroke, epilepsy, dystonia, noncompaction, and kidney agenesis requires genetic work-up. <i>Child's Nervous System</i> , 2020, 36, 1831-1832.	0.6	0
232	Heterogeneous phenotypic expression of C1QBP variants is attributable to variable heteroplasmy of secondary mtDNA deletions and mtDNA copy number. <i>Human Mutation</i> , 2020, 41, 2012-2013.	1.1	1
233	Stroke-like Episodes in m.3243A>G Carriers Need to Be Monitored by MRI Starting with the Onset of Clinical Manifestations. <i>American Journal of Neuroradiology</i> , 2020, 41, E17-E18.	1.2	0
234	Mitochondrial disorder should be considered as a differential of late-onset myasthenia gravis. <i>Acta Neurologica Belgica</i> , 2020, 121, 1891-1892.	0.5	0

#	ARTICLE	IF	CITATIONS
235	Is the spinal cord truly affected in half of the patients with Kearns-Sayre syndrome?. <i>Neuroradiology</i> , 2020, 62, 1205-1206.	1.1	1
236	Aphatic epileptic state as a manifestation of a stroke-like episode in MELAS. <i>Epilepsy Research</i> , 2020, 180, 106531.	0.8	1
237	Update on the neurology of COVID-19. <i>Journal of Medical Virology</i> , 2020, 92, 2316-2318.	2.5	32
238	Multisystem Myotilinopathy, including Myopathy and Left Ventricular Noncompaction, due to the MYOT Variant c.179C>T. <i>Case Reports in Cardiology</i> , 2020, 2020, 1-4.	0.1	2
239	Recognise subtle manifestations for classifying MELAS. <i>Journal of the Neurological Sciences</i> , 2020, 414, 116926.	0.3	1
240	Systemic toxicity to betamethasone ointment. <i>Clinical Case Reports (discontinued)</i> , 2020, 8, 1635-1637.	0.2	1
241	We never speak about sudden unexpected death in Parkinson's disease. <i>European Journal of Neurology</i> , 2020, 27, e30.	1.7	1
242	Unusual Phenotype and Disease Trajectory in Kearns-Sayre Syndrome. <i>Case Reports in Neurological Medicine</i> , 2020, 2020, 1-6.	0.3	5
243	Monogenic cerebral small-vessel diseases: diagnosis and therapy. Consensus recommendations of the European Academy of Neurology. <i>European Journal of Neurology</i> , 2020, 27, 909-927.	1.7	103
244	Mitochondrial disorders are prone to propofol infusion syndrome. <i>Acute Medicine & Surgery</i> , 2020, 7, e495.	0.5	0
245	Clinical Therapeutic Management of Human Mitochondrial Disorders. <i>Pediatric Neurology</i> , 2020, 113, 66-74.	1.0	6
246	Early white matter changes on diffusion tensor imaging in amyotrophic lateral sclerosis. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 1265-1265.	1.7	1
247	Double Trouble from POLG1 and CLCN1 Variants with Intrafamilial Phenotypic Heterogeneity. <i>Movement Disorders Clinical Practice</i> , 2020, 7, 575-576.	0.8	0
248	Comment on Progression of Retinopathy Secondary to Maternally Inherited Diabetes and Deafness: Evaluation of Predicting Parameters. <i>American Journal of Ophthalmology</i> , 2020, 216, 283-284.	1.7	0
249	Metabolic stroke or stroke-like lesion: Peculiarities of a phenomenon. <i>Journal of the Neurological Sciences</i> , 2020, 412, 116726.	0.3	51
250	Incapacitating, generalised myalgias and muscle stiffness under duloxetine and aripiprazole. <i>International Journal of Clinical Practice</i> , 2020, 74, e13487.	0.8	1
251	<i>POLG1</i>-related phenotypes are heterogeneous and progressive due to secondary mtDNA maintenance defects. <i>International Journal of Neuroscience</i> , 2020, 130, 1282-1283.	0.8	0
252	Variant m.1555A>G in MT-RNR1 causes hearing loss and multiorgan mitochondrial disorder. <i>Medicine (United States)</i> , 2020, 99, e18488.	0.4	2

#	ARTICLE	IF	CITATIONS
253	Re: Parisi et Al.: Functional changes of retinal ganglion cells and visual pathways in patients with chronic Leber's hereditary optic neuropathy during one year of follow-up (Ophthalmology.) Tj ETQq1 1 0.7843 14rgBT /Overlock 10		
254	Assessing the effect of non-invasive ventilation on cardiac function in Duchenne muscular dystrophy requires prospective studies. <i>Cardiology in the Young</i> , 2020, 30, 1215-1216.	0.4	1
255	Initial deterioration upon intravenous methyl-prednisolon in myasthenia is multifactorial. <i>Journal of the Neurological Sciences</i> , 2020, 412, 116812.	0.3	0
256	Assessment of the Idebenone Effect on LHON Eyes Requires High-quality Studies. <i>Current Eye Research</i> , 2020, 45, 1451-1452.	0.7	0
257	Factors determining the early beneficial effect of unilateral gene therapy in Leber's hereditary optic neuropathy patients carrying variant m.11778G>A. <i>Acta Ophthalmologica</i> , 2020, 98, e788.	0.6	0
258	Comment on "CPEO and Mitochondrial Myopathy in a Patient with DGUOK Compound Heterozygous Pathogenetic Variant and mtDNA Multiple Deletions": Case Reports in Neurological Medicine, 2020, 2020, 1-2.	0.3	0
259	Update Review about Metabolic Myopathies. <i>Life</i> , 2020, 10, 43.	1.1	11
260	Causes of hypogeusia/hyposmia in SARS-CoV2 infected patients. <i>Journal of Medical Virology</i> , 2020, 92, 1793-1794.	2.5	92
261	Primary hypoparathyroidism and multiple neuraxial involvement in mitochondrial disorder due to the variant m.15043G>A in MT-CYB. <i>Journal of the Neurological Sciences</i> , 2020, 414, 116853.	0.3	3
262	SARS-CoV-2-Associated Acute Hemorrhagic, Necrotizing Encephalitis (AHNE) Presenting with Cognitive Impairment in a 44-Year-Old Woman without Comorbidities: A Case Report. <i>American Journal of Case Reports</i> , 2020, 21, e925641.	0.3	54
263	Phenotypic Heterogeneity in 5 Family Members with the Mitochondrial Variant m.3243A>G. <i>American Journal of Case Reports</i> , 2020, 21, e927938.	0.3	8
264	The metabolic hypothesis is more likely than the epileptogenic hypothesis to explain stroke-like lesions. <i>Wellcome Open Research</i> , 2020, 5, 51.	0.9	5
265	The mitochondrial calcium uniporter: a new therapeutic target for Parkinson's disease-related cardiac dysfunctions?. <i>Clinics</i> , 2020, 75, e1299.	0.6	8
266	Alcohol and sudden unexpected death in epilepsy: do not pop the cork. <i>Clinics</i> , 2020, 75, e1770.	0.6	5
267	Atherosclerosis Can Be Mitochondrial: A Review. <i>Cureus</i> , 2020, 12, e6987.	0.2	7
268	Letter to the Editor: Comments on "Obesity associated with a novel mitochondrial tRNACys m.5802A>G mutation in a Chinese family": <i>Bioscience Reports</i> , 2020, 40, .	1.1	0
269	Stroke in MELAS is a Vasogenic Edema and not Ischemic. <i>Basic and Clinical Neuroscience</i> , 2020, 11, 379-380.	0.3	0
270	Duchenne muscular dystrophy requires treatment also of cardiac, respiratory, cerebral, and orthopedic compromise. <i>Arquivos De Neuro-Psiquiatria</i> , 2020, 78, 313-314.	0.3	0

#	ARTICLE	IF	CITATIONS
271	Considerations on the management of pyruvate dehydrogenase deficiency. <i>Folia Morphologica</i> , 2020, 79, 415-416.	0.4	0
272	Detection of compound heterozygous variants in LPIN1 does not necessarily imply pathogenicity in a patient with rhabdomyolysis. <i>F1000Research</i> , 2020, 9, 15.	0.8	0
273	Rare Phenotypic Manifestations of MELAS. <i>Yonsei Medical Journal</i> , 2020, 61, 904.	0.9	3
274	The Leigh phenotype resulting from C12orf65 variants. <i>Genetics and Molecular Biology</i> , 2020, 43, e20200177.	0.6	1
275	Cluster headache as a manifestation of a stroke-like episode in a carrier of the <i>MT-ND3</i> variant m.10158T>C. <i>World Journal of Clinical Cases</i> , 2020, 8, 242-244.	0.3	0
276	The metabolic hypothesis is more likely than the epileptogenic hypothesis to explain stroke-like lesions. <i>Wellcome Open Research</i> , 2020, 5, 51.	0.9	3
277	Uncovering the etiology of ptosis prior to blepharoplasty. <i>Archives of Plastic Surgery</i> , 2020, 47, 487-487.	0.4	0
278	Phenotypic spectrum of variants in the beta-oxidation enoyl-CoA hydratase-1 (ECHS-1) gene. <i>European Journal of Paediatric Neurology</i> , 2020, 29, 101-102.	0.7	0
279	Comment on: Sideroblastic anemia associated with multisystem mitochondrial disorders: The phenotypic spectrum of <i>PUS1</i> and <i>COX10</i> variants and mtDNA deletions needs to be prospectively assessed. <i>Pediatric Blood and Cancer</i> , 2019, 66, e27945.	0.8	1
280	Extensive clinical and genetic workup is worthwhile in patients with Leigh-like syndrome due to the TSFM variant c.547G>A. <i>Neurogenetics</i> , 2019, 20, 219-220.	0.7	0
281	Elucidation of the genetic background in familial multiple sclerosis requires genetic work-up. <i>Clinical Neurology and Neurosurgery</i> , 2019, 184, 105425.	0.6	1
282	Comment on "Proximal Myopathy due to m.5835G>A Mutation in Mitochondrial MT-TY Gene": Case Reports in <i>Neurological Medicine</i> , 2019, 2019, 1-2.	0.3	0
283	Noncompaction and Takotsubo Syndrome in a Neuromuscular Disorder. <i>Case Reports in Cardiology</i> , 2019, 2019, 1-3.	0.1	1
284	Cerebellar Stroke-like Lesions?. <i>Internal Medicine</i> , 2019, 58, 3205-3205.	0.3	3
285	Commentary: New Variant of MELAS Syndrome With Executive Dysfunction, Heteroplasmic Point Mutation in the MT-ND4 Gene (m.12015T>C; p.Leu419Pro) and Comorbid Polyglandular Autoimmune Syndrome Type 2. <i>Frontiers in Immunology</i> , 2019, 10, 1333.	2.2	1
286	Cerebral imaging in adult mitochondrial disorders. <i>Journal of the Neurological Sciences</i> , 2019, 404, 29-35.	0.3	5
287	Central nervous system abnormalities in spinal and bulbar muscular atrophy (Kennedy's disease). <i>Clinical Neurology and Neurosurgery</i> , 2019, 184, 105426.	0.6	6
288	<p>Barth syndrome: mechanisms and management</p>. <i>The Application of Clinical Genetics</i> , 2019, Volume 12, 95-106.	1.4	30

#	ARTICLE	IF	CITATIONS
289	<p></p>Ketogenic diet in ATAD3A mutation carriers may not improve cerebellar atrophy but some clinical features [Letter]. The Application of Clinical Genetics, 2019, Volume 12, 161-162.	1.4	1
290	Commentary: Epilepsy in Leigh Syndrome With Mitochondrial DNA Mutations. Frontiers in Neurology, 2019, 10, 973.	1.1	1
291	Only pathogenic variants in protein-coding mtDNA genes cause Leigh syndrome. Journal of the Neurological Sciences, 2019, 407, 116447.	0.3	0
292	Reply to: "Advances in imaging of brain abnormalities in neuromuscular disease". Therapeutic Advances in Neurological Disorders, 2019, 12, 175628641987832.	1.5	0
293	Neuropathy due to impaired axonal transport of non-fragmented mitochondria in MYH14 mutation carriers. EBioMedicine, 2019, 49, 24.	2.7	1
294	REM sleep without atonia as prodromal marker of Lewy body disease: Fake news or the real deal? Parkinsonism and Related Disorders, 2019, 67, 34-35.	1.1	2
295	Sudden unexpected death in Parkinson's disease: why is drinking water important?. Neurodegenerative Disease Management, 2019, 9, 241-246.	1.2	7
296	Simultaneous transient global amnesia and Takotsubo syndrome after death of a relative: a case report. Journal of Medical Case Reports, 2019, 13, 22.	0.4	7
297	PTCD3 mutations cause Leigh-like rather than Leigh syndrome. Neurogenetics, 2019, 20, 53-54.	0.7	1
298	Liability of sepsis is hardly determined by the <i>COXI</i> variant m.6459T>C. Journal of Cellular and Molecular Medicine, 2019, 23, 689-690.	1.6	1
299	Evidence for laminar cortical necrosis as histological equivalent of a non-classical stroke-like lesion in an m.8344A>G carrier. Neuropathology, 2019, 39, 324-325.	0.7	1
300	Folinic acid is ineffective for treating Kearns-Sayre syndrome. Neurologia (English Edition), 2019, 34, 347.	0.2	0
301	Comment on APLAR Grand Round: Phenotypes of single mtDNA deletions may unequivocally suggest mitochondrial disease. International Journal of Rheumatic Diseases, 2019, 22, 1165-1166.	0.9	0
302	Autosomal dominant optic atrophy plus due to the novel OPA1 variant c.1463G>C. Metabolic Brain Disease, 2019, 34, 1023-1027.	1.4	2
303	Pharmacotherapeutic management of epilepsy in MERRF syndrome. Expert Opinion on Pharmacotherapy, 2019, 20, 1289-1297.	0.9	8
304	Diagnostic and therapeutic aspects of Leigh syndrome due to the variant m.10197G>A. Journal of the Neurological Sciences, 2019, 400, 182-183.	0.3	1
305	Before attributing CPEO and ptosis to the variant m.14819T>G its pathogenicity needs to be established. Journal of the Neurological Sciences, 2019, 401, 110-111.	0.3	0
306	Methyl-DOPA causing reversible peripheral facial palsy. Clinical Neurology and Neurosurgery, 2019, 179, 53-54.	0.6	1

#	ARTICLE	IF	CITATIONS
307	Mitochondrial metabolic stroke: Phenotype and genetics of stroke-like episodes. <i>Journal of the Neurological Sciences</i> , 2019, 400, 135-141.	0.3	50
308	Comments on: Isolated Noncompaction Suggests Subclinical Myopathy. <i>Revista Brasileira De Ginecologia E Obstetricia</i> , 2019, 41, 135-136.	0.3	0
309	Peculiarities of stroke-like lesions on MRI. <i>European Journal of Radiology Open</i> , 2019, 6, 60-61.	0.7	3
310	Re: Increased incidence of obstetric complications in women carrying mitochondrial DNA mutations: a retrospective cohort study in a single tertiary centre. <i>BJOG: an International Journal of Obstetrics and Gynaecology</i> , 2019, 126, 1400-1401.	1.1	1
311	Genetic Data Are a Prerequisite for Interpreting Clinical and Muscle Biopsy Findings in MELAS. <i>Yonsei Medical Journal</i> , 2019, 60, 399.	0.9	1
312	Early-age Ndufs4 knockout mice are an inappropriate animal model of Leigh syndrome. <i>Radiological Physics and Technology</i> , 2019, 12, 230-231.	1.0	1
313	Mitochondrial Ophthalmoplegia Is Not Only due to mtDNA Deletions. <i>Yonsei Medical Journal</i> , 2019, 60, 230.	0.9	0
314	Whole exome should be preferred over Sanger sequencing in suspected mitochondrial myopathy. <i>Neurobiology of Aging</i> , 2019, 78, 166-167.	1.5	0
315	Heteroplasmy Rates of the m.14495A>G variant in MT-ND6 May Not Predict the Phenotype of LHON. <i>Translational Vision Science and Technology</i> , 2019, 8, 42.	1.1	0
316	Mitochondrial Parkinsonism due to <i>SPG7/Paraplegin</i> variants with secondary mtDNA depletion. <i>Movement Disorders</i> , 2019, 34, 1931-1932.	2.2	0
317	Commentary: Takotsubo Cardiomyopathy-Acute Cardiac Dysfunction Associated With Neurological and Psychiatric Disorders. <i>Frontiers in Neurology</i> , 2019, 10, 1163.	1.1	3
318	Multisystem mitochondrial disorder is more prevalent than BGC1 variants in patients with Fahr's syndrome. <i>ENeurologicalSci</i> , 2019, 17, 100216.	0.5	0
319	MELAS in a Walk-in Customer. <i>Journal of Neurosciences in Rural Practice</i> , 2019, 10, 725-727.	0.3	0
320	The "Toenail Sign" in MELAS May Be the End Stage of a Stroke-like Lesion. <i>Pediatric Neurology</i> , 2019, 93, 67.	1.0	0
321	Survival and outcome in MELAS not only depends on onset and disease duration. <i>Journal of the Neurological Sciences</i> , 2019, 397, 9-10.	0.3	0
322	Maternal transmission of CNTN6 copy number variation suggests mitochondrial disorder. <i>Schizophrenia Research</i> , 2019, 206, 454-455.	1.1	0
323	Antiepileptics and NO-precursors may be beneficial for stroke-like episodes. <i>ENeurologicalSci</i> , 2019, 14, 38-39.	0.5	1
324	Management of NARS2-Related Mitochondrial Disorder is Complex. <i>Pediatric Neurology</i> , 2019, 93, 64.	1.0	0

#	ARTICLE	IF	CITATIONS
325	Sudden unexpected death in epilepsy: Rethinking the unthinkable. <i>Epilepsy and Behavior</i> , 2019, 93, 148-149.	0.9	6
326	Tinnitus is multicausal and may not only be related to DNA variants. <i>European Archives of Oto-Rhino-Laryngology</i> , 2019, 276, 1551-1552.	0.8	0
327	Silencing of FOXRED1 in C57b1/6 mice does not generate an appropriate animal model of Leigh syndrome. <i>Metabolic Brain Disease</i> , 2019, 34, 1-2.	1.4	3
328	Hereditary transthyretin-related amyloidosis. <i>Acta Neurologica Scandinavica</i> , 2019, 139, 92-105.	1.0	35
329	Patients with MELAS not only require treatment of stroke-like episodes but a comprehensive individual and family management. <i>Acta Neurologica Scandinavica</i> , 2019, 139, 86-87.	1.0	1
330	Mitochondrial disorder mimicking rheumatoid disease. <i>Zeitschrift Fur Rheumatologie</i> , 2019, 78, 875-880.	0.5	8
331	Sudden unexpected death in Parkinson's disease: why do neuroscientists still ignore this condition?. <i>Neurological Sciences</i> , 2019, 40, 413-414.	0.9	2
332	El Ácido foliÁnico no es eficaz en el tratamiento del sÁndrome de Kearns-Sayre. <i>NeurologÁa</i> , 2019, 34, 347.	0.3	0
333	ADVERSE REACTION TO ANESTHESIA IN A M.8993T>C CARRIER WITH LEIGH SYNDROME. <i>Revista Paulista De Pediatria</i> , 2019, 37, 135-136.	0.4	0
334	Doping May Be Responsible for De Novo Mitochondrial Disorder. <i>Advanced Pharmaceutical Bulletin</i> , 2019, 9, 180-181.	0.6	0
335	Genetic work-up of hereditary spastic paraplegias is crucial for classifying these disorders. <i>Arquivos De Neuro-Psiquiatria</i> , 2019, 77, 597-597.	0.3	1
336	Retinal nerve fiber and ganglion cell layer thinning in hereditary and acquired mitochondrial optic neuropathies. <i>International Journal of Ophthalmology</i> , 2019, 12, 1666-1666.	0.5	0
337	MELAS Missed for Years: Stroke-Like Lesions Are No Indication for Brain Biopsy. <i>Case Reports in Neurological Medicine</i> , 2019, 2019, 1-4.	0.3	0
338	Early identification of LHON carriers may improve outcome. <i>Romanian Journal of Ophthalmology</i> , 2019, 63, 102-103.	0.4	0
339	LHON needs to be genetically diagnosed and the idebenone effect quantified. <i>Romanian Journal of Ophthalmology</i> , 2019, 63, 306-307.	0.4	1
340	Headache in mitochondrial disorders. <i>Clinical Neurology and Neurosurgery</i> , 2018, 166, 44-49.	0.6	14
341	Ketogenic diet and avoidance of mitochondrion-toxic AEDs may improve the outcome of mitochondrial epilepsy. <i>Clinical Neurology and Neurosurgery</i> , 2018, 173, 202-203.	0.6	2
342	Re: Guy etÁal.: Gene therapy for Leber hereditary optic neuropathy: low-and medium-dose visual results (<i>Ophthalmology</i> . 2017;124:1621-1634). <i>Ophthalmology</i> , 2018, 125, e14-e15.	2.5	5

#	ARTICLE	IF	CITATIONS
343	Only some patients with bulbar and spinal muscular atrophy may develop cardiac disease. <i>Molecular Genetics and Metabolism Reports</i> , 2018, 14, 44-46.	0.4	2
344	Phenotypic heterogeneity of <i>POMT2</i> gene variants. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 743-745.	0.7	1
345	Coronary ectasia in amyloid cardiomyopathy and neuropathy due to the transthyretin mutation c.323A>G. <i>Heart and Lung: Journal of Acute and Critical Care</i> , 2018, 47, 127-129.	0.8	1
346	MERRF Classification: Implications for Diagnosis and Clinical Trials. <i>Pediatric Neurology</i> , 2018, 80, 8-23.	1.0	58
347	Cerebellar atrophy is common among mitochondrial disorders. <i>Metabolic Brain Disease</i> , 2018, 33, 987-988.	1.4	6
348	Cardiac abnormalities in Parkinsonâ€™s disease and Parkinsonism. <i>Journal of Clinical Neuroscience</i> , 2018, 53, 1-5.	0.8	100
349	Sudden Unexpected Death in Parkinsonâ€™s Disease (SUDPAR): a fatal event that James Parkinson did not address. <i>Age and Ageing</i> , 2018, 47, 627-627.	0.7	2
350	Neuropathy of peripheral nerves in Leber's hereditary optic neuropathy. <i>Journal of the Neurological Sciences</i> , 2018, 390, 193-194.	0.3	1
351	Wolff-Parkinson-White syndrome and noncompaction in Leberâ€™s hereditary optic neuropathy due to the variant m.3460G>A. <i>Journal of International Medical Research</i> , 2018, 46, 2054-2060.	0.4	9
352	Polymegathism as a biomarker of mitochondrial disorders. <i>Graefe's Archive for Clinical and Experimental Ophthalmology</i> , 2018, 256, 1211-1212.	1.0	0
353	Muscle, cardiac, and cerebral manifestations in female carriers of dystrophin variants. <i>Journal of the Neurological Sciences</i> , 2018, 388, 107-108.	0.3	8
354	Mitochondrial disorders of the retinal ganglion cells and the optic nerve. <i>Mitochondrion</i> , 2018, 42, 1-10.	1.6	18
355	The Cerebellum Is a Common Site of Affection in Leigh Syndrome. <i>Pediatric Neurology</i> , 2018, 78, e9.	1.0	1
356	Phenotypic spectrum of POLG1 mutations. <i>Neurological Sciences</i> , 2018, 39, 571-573.	0.9	6
357	The Tip of the Iceberg in Maternally Inherited Diabetes and Deafness. <i>Oman Medical Journal</i> , 2018, 33, 437-440.	0.3	3
358	Involvement of the Spinal Cord in Mitochondrial Disorders. <i>Journal of Neurosciences in Rural Practice</i> , 2018, 09, 245-251.	0.3	23
359	Retinal involvement in m.3243A>G carriers. <i>Ophthalmic Genetics</i> , 2018, 39, 664-665.	0.5	1
360	An update on diagnosis and therapy of metabolic myopathies. <i>Expert Review of Neurotherapeutics</i> , 2018, 18, 933-943.	1.4	7

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361	Phenytoin-induced choreoathetosis after serial seizures due to traumatic brain injury and chronic alcoholism. <i>Clinical Case Reports (discontinued)</i> , 2018, 6, 2316-2318.	0.2	3
362	Letter to the Editor: Endocrine Compromise in Mitochondrial Disorders. <i>Journal of the Endocrine Society</i> , 2018, 2, 570-571.	0.1	3
363	Genotypic and Phenotypic Heterogeneity of LGMD1D due to <i>DNAJB6</i> Mutations. <i>Yonsei Medical Journal</i> , 2018, 59, 1008.	0.9	2
364	Aortic root ectasia as a phenotypic feature of a mitochondrial disorder. <i>Clinical Case Reports (discontinued)</i> , 2018, 6, 1501-1503.	0.2	1
365	Kearns-Sayre syndrome is genetically and phenotypically heterogeneous. <i>Pediatrica Medica E Chirurgica</i> , 2018, 40, .	0.1	4
366	Concomitant cardiac and cerebral takotsubo syndrome requires specific management. <i>Clinical Autonomic Research</i> , 2018, 28, 601-602.	1.4	0
367	MELAS can be psychiatric and neurological. <i>ENeurologicalSci</i> , 2018, 11, 3-4.	0.5	2
368	Infantile-onset deafness in m.7445A>G carriers may be multicausal. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2018, 111, 192-193.	0.4	1
369	Comment on "Role of Mitochondrial Genome Mutations in Pathogenesis of Carotid Atherosclerosis". <i>Oxidative Medicine and Cellular Longevity</i> , 2018, 2018, 1-2.	1.9	1
370	The Genetic Cause of Kearns-Sayre Syndrome Determines Counselling and Outcome of These Patients. <i>Canadian Journal of Cardiology</i> , 2018, 34, 1234.e7.	0.8	3
371	Takotsubo as Initial Manifestation of Non-Myopathic Cardiomyopathy Due to the Titin Variant c.1489G>T. <i>Medicines (Basel, Switzerland)</i> , 2018, 5, 80.	0.7	4
372	Features on cerebral imaging suggesting mitochondrial disorder. <i>Neurodegenerative Disease Management</i> , 2018, 8, 215-216.	1.2	3
373	Cerebral imaging in paediatric mitochondrial disorders. <i>Neuroradiology Journal</i> , 2018, 31, 596-608.	0.6	8
374	Biomarkers for Detecting Mitochondrial Disorders. <i>Journal of Clinical Medicine</i> , 2018, 7, 16.	1.0	30
375	Mitochondrial ataxia is genetically and phenotypically heterogeneous. <i>CNS Neuroscience and Therapeutics</i> , 2018, 24, 1301-1302.	1.9	0
376	Modified Yarham and Smith scores for pathogenicity assessment of mtDNA tRNA variants. <i>Neuromuscular Disorders</i> , 2018, 28, 373-374.	0.3	0
377	Insular cortex lesions are not the only culprit in Takotsubo syndrome. <i>Clinical Cardiology</i> , 2018, 41, 1407-1408.	0.7	2
378	Respiratory insufficiency from myasthenia gravis and polymyositis due to malignant thymoma triggering Takotsubo syndrome. <i>International Journal of Neuroscience</i> , 2018, 128, 1207-1210.	0.8	7

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379	Gastrointestinal Involvement in m.3243A>G-associated MELAS. <i>Internal Medicine</i> , 2018, 57, 769-770.	0.3	5
380	Clinical Perspectives of Mitochondrial Disorders. <i>Pediatric Endocrinology Reviews</i> , 2018, 16, 203-208.	1.2	7
381	Myotonic dystrophy-2: Unusual phenotype due to a small CCTG-expansion. <i>Balkan Journal of Medical Genetics</i> , 2018, 21, 39-43.	0.5	2
382	Levels of nitric oxide pathway parameters may depend on heteroplasmy rates of the m.3243A>G mutation. <i>International Journal of Cardiology</i> , 2017, 229, 26.	0.8	1
383	Left ventricular noncompaction cardiomyopathy: cardiac, neuromuscular, and genetic factors. <i>Nature Reviews Cardiology</i> , 2017, 14, 224-237.	6.1	166
384	Growth-hormone deficiency in mitochondrial disorders. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2017, 30, 479-481.	0.4	3
385	Low blood heteroplasmy-rate may cause late-onset MELAS. <i>Molecular Genetics and Metabolism Reports</i> , 2017, 10, 100.	0.4	1
386	Takotsubo in Miller-Fisher and Guillain Barre syndrome. <i>American Journal of Emergency Medicine</i> , 2017, 35, 1201-1202.	0.7	1
387	Death in Pediatric Mitochondrial Disorders. <i>Pediatric Neurology</i> , 2017, 73, e1.	1.0	2
388	Transient global amnesia: The cerebral Takotsubo?. <i>Journal of the Neurological Sciences</i> , 2017, 376, 196-197.	0.3	15
389	Renal manifestations of primary mitochondrial disorders. <i>Biomedical Reports</i> , 2017, 6, 487-494.	0.9	57
390	Is the Takotsubo syndrome a brain-heart or multiorgan disorder?. <i>Journal of the Neurological Sciences</i> , 2017, 378, 239-240.	0.3	1
391	Unilateral Ptosis and Homolateral Hemifacial Weakness in Chronic Progressive External Ophthalmoplegia. <i>Neuro-Ophthalmology</i> , 2017, 41, 165-166.	0.4	1
392	Epilepsy in MELAS. <i>Pediatric Neurology</i> , 2017, 67, e7-e8.	1.0	1
393	Takotsubo in amyotrophic lateral sclerosis. <i>Journal of the Neurological Sciences</i> , 2017, 379, 339-340.	0.3	1
394	Genetic Counselling for Maternally Inherited Mitochondrial Disorders. <i>Molecular Diagnosis and Therapy</i> , 2017, 21, 419-429.	1.6	93
395	NDUFS4-related Leigh syndrome in Hutterites. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 1450-1451.	0.7	5
396	Gastrointestinal manifestations of mitochondrial disorders: a systematic review. <i>Therapeutic Advances in Gastroenterology</i> , 2017, 10, 142-154.	1.4	63

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397	Causally treatable, hereditary neuropathies in Fabry's disease, transthyretin-related familial amyloidosis, and Pompe's disease. <i>Acta Neurologica Scandinavica</i> , 2017, 136, 558-569.	1.0	10
398	Onset of MELAS due to the m.3243A > G mutation is early if the large phenotypic variability is considered. <i>Molecular Genetics and Metabolism Reports</i> , 2017, 10, 23.	0.4	2
399	Phenotypic heterogeneity of MELAS. <i>Molecular Genetics and Metabolism Reports</i> , 2017, 10, 18-19.	0.4	4
400	Pathogenicity of the LHON variant m.3472T > C is uncertain. <i>Journal of the Neurological Sciences</i> , 2017, 382, 164-165.	0.3	0
401	Cerebral Manifestations of Mitochondrial Disorders. <i>Canadian Journal of Neurological Sciences</i> , 2017, 44, 654-663.	0.3	6
402	Can <sc>MR</sc> spectroscopy and muscle biopsy findings be correlated with <sc>MELAS</sc> and <sc>CPEO</sc>?. <i>CNS Neuroscience and Therapeutics</i> , 2017, 23, 846-847.	1.9	0
403	Affection of immune cells by a C10orf2 mutation manifesting as mitochondrial myopathy and transient sensory transverse syndrome. <i>Acta Neurologica Belgica</i> , 2017, 117, 969-970.	0.5	6
404	Compromiso renal en MELAS. <i>Medicina Clínica</i> , 2017, 149, 314.	0.3	1
405	Mitochondrial cardioencephalopathy due to a COQ4 mutation. <i>Molecular Genetics and Metabolism Reports</i> , 2017, 13, 7-8.	0.4	1
406	Kearns-Sayre syndrome in the absence of a mtDNA deletion?. <i>Andrologia</i> , 2017, 49, e12810.	1.0	0
407	Management of epilepsy in MERRF syndrome. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2017, 50, 166-170.	0.9	22
408	Do lesional perfusion abnormalities on arterial spin labeling truly contribute to the diagnosis of Leigh syndrome?. <i>Pediatric Radiology</i> , 2017, 47, 124-125.	1.1	1
409	Why does Leigh syndrome respond to immunotherapy?. <i>Molecular Genetics and Metabolism Reports</i> , 2017, 11, 90-91.	0.4	3
410	Ophthalmologic involvement in mitochondrial disorders. <i>Ophthalmic Genetics</i> , 2017, 38, 298-298.	0.5	2
411	Mitochondrial multiorgan disorder syndrome score generated from definite mitochondrial disorders. <i>Neuropsychiatric Disease and Treatment</i> , 2017, Volume 13, 2569-2579.	1.0	18
412	Psychological morbidity in Leber's hereditary optic neuropathy depends on phenotypic, social, economic, and genetic factors. <i>Clinical Ophthalmology</i> , 2017, Volume 11, 959-962.	0.9	1
413	Sudden unexpected death in Parkinson's disease (SUDPAR): a review of publications since the decade of the brain. <i>Clinics</i> , 2017, 72, 649-651.	0.6	41
414	Noncompaction may not only be non-isolated but also myopathic. Commentary to the article: "Left ventricular non-compaction associated with hypertrophic cardiomyopathy in the same patient". <i>Kardiologia Polska</i> , 2017, 75, 726-726.	0.3	0

#	ARTICLE	IF	CITATIONS
415	Psoriasis, bulbar involvement, and diarrhea in late myoclonic epilepsy with ragged-red fibers-syndrome due to the m.8344A > G tRNA (Lys) mutation. Iranian Journal of Neurology, 2017, 16, 45-49.	0.5	5
416	Diagnosis of Kearns-Sayre Syndrome Requires Comprehensive Work-up. Chinese Medical Journal, 2016, 129, 2518-2519.	0.9	1
417	Leber’s hereditary optic neuropathy is multiorgan not mono-organ. Clinical Ophthalmology, 2016, Volume 10, 2187-2190.	0.9	40
418	Comment on "Symptomatic Trifascicular Block in Steinert's Disease: Is It Too Soon for a Pacemaker?" Case Reports in Cardiology, 2016, 2016, 1-2.	0.1	1
419	Noncompaction and Dilated Cardiomyopathy in a Patient with Schizophrenia. Case Reports in Cardiology, 2016, 2016, 1-3.	0.1	1
420	Comment on "A Rare Case of Renal Infarct due to Noncompaction Cardiomyopathy: A Case Report and Literature Review" Case Reports in Cardiology, 2016, 2016, 1-2.	0.1	0
421	Diagnose Kearns's Sayre syndrome genetically and investigate the phenotype comprehensively. Oxford Medical Case Reports, 2016, 2016, omw059.	0.2	1
422	Mitochondrial vasculopathy. World Journal of Cardiology, 2016, 8, 333.	0.5	51
423	Secondary myopathy due to systemic diseases. Acta Neurologica Scandinavica, 2016, 134, 388-402.	1.0	17
424	Peculiarities of progressive external ophthalmoplegia due to single mitochondrial DNA deletions. Journal of the Formosan Medical Association, 2016, 115, 1099-1100.	0.8	2
425	Prevalence of neoplasms in definite and probable mitochondrial disorders. Mitochondrion, 2016, 29, 31-34.	1.6	21
426	Stress from myasthenic crisis triggers Takotsubo (broken heart) syndrome. International Journal of Cardiology, 2016, 203, 616-617.	0.8	16
427	Toxicity of Antiepileptic Drugs to Mitochondria. Handbook of Experimental Pharmacology, 2016, 240, 473-488.	0.9	65
428	Nephrocalcinosis and retinal dystrophy, rare manifestations of MPV17-related mitochondrial depletion syndrome?. Molecular Genetics and Metabolism Reports, 2016, 9, 18.	0.4	1
429	Mitochondrial vasculopathy due to the m.3243A>G mutation is not restricted to the carotid artery. Molecular Genetics and Metabolism Reports, 2016, 9, 34.	0.4	1
430	Motor neuron disease triggering takotsubo syndrome. International Journal of Cardiology, 2016, 223, 21-22.	0.8	3
431	Mitochondrial movement disorders. Revue Neurologique, 2016, 172, 716-717.	0.6	1
432	Cerebral involvement in mitochondrial disorders on imaging. Child's Nervous System, 2016, 32, 2059-2060.	0.6	3

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433	Hearing impairment in m.3243A>G carriers requires comprehensive work- and follow-up. <i>Clinical Neurology and Neurosurgery</i> , 2016, 150, 199-200.	0.6	1
434	Re: Feuer et al.: Gene therapy for Leber hereditary optic neuropathy: initial results (<i>Ophthalmology</i>) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5	2.5	2
435	Cardiac disease in brain-heart disorders. <i>Acta Cardiologica</i> , 2016, 71, 389-394.	0.3	3
436	Glucocorticoids for treating Takotsubo syndrome?. <i>Clinical Autonomic Research</i> , 2016, 26, 159-159.	1.4	0
437	Phenotypic spectrum of the m.8344A>G mutation. <i>Journal of Neurology</i> , 2016, 263, 1452-1453.	1.8	1
438	Wet, volatile, and dry biomarkers of exercise-induced muscle fatigue. <i>BMC Musculoskeletal Disorders</i> , 2016, 17, 40.	0.8	25
439	Lactose-free diet inducing aseptic pancreatitis and myoclonic jerks in late-onset, putative MERRF syndrome. <i>Journal of Neurology</i> , 2016, 263, 586-587.	1.8	3
440	Oncocytoma and noncompaction in metabolic myopathy. <i>International Journal of Cardiology</i> , 2016, 207, 382-383.	0.8	4
441	The Eye on Mitochondrial Disorders. <i>Journal of Child Neurology</i> , 2016, 31, 652-662.	0.7	23
442	Cardiac disease in brain-heart disorders. <i>Acta Cardiologica</i> , 2016, 71, 389-94.	0.3	1
443	Malingering and Factitious Disorder (Munchausensyndrome) can be Mitochondrial. <i>Indian Journal of Psychological Medicine</i> , 2016, 38, 348-352.	0.6	2
444	Considerations about the genetics of left ventricular hypertrabeculation/non-compaction. <i>Cardiology in the Young</i> , 2015, 25, 1435-1437.	0.4	17
445	Multiorgan disorder syndrome (MODS) in an octagenarian suggests mitochondrial disorder. <i>Revista Medica De Chile</i> , 2015, 143, 1210-1214.	0.1	20
446	Diabetes in Kearns-Sayre Syndrome: More Common than Anticipated. <i>Canadian Journal of Diabetes</i> , 2015, 39, 253.	0.4	6
447	Therapeutic Strategies for Mitochondrial Disorders. <i>Pediatric Neurology</i> , 2015, 52, 302-313.	1.0	34
448	Leriche-syndrome despite regular sport and non-compaction suggest neuromuscular disease. <i>International Journal of Cardiology</i> , 2015, 191, 15-17.	0.8	4
449	Noncompaction with dysmorphism, mental retardation, general wasting, and hypogonadism requires neurologic and sophisticated cytogenetic investigations. <i>Anatolian Journal of Cardiology</i> , 2015, 15, 433-434.	0.5	0
450	Age-Dependency of Cardiac and Neuromuscular Findings in Adults With Left Ventricular Hypertrabeculation/Noncompaction. <i>American Journal of Cardiology</i> , 2015, 115, 1287-1292.	0.7	17

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451	Exhaustion or fatigability may not only be cardiac but also myopathic. <i>Netherlands Heart Journal</i> , 2015, 23, 292-293.	0.3	0
452	Abnormalities of Skin and Cutaneous Appendages in Neuromuscular Disorders. <i>Pediatric Neurology</i> , 2015, 53, 301-308.	1.0	5
453	Glucocorticoids for mitochondrial disorders. <i>Singapore Medical Journal</i> , 2015, 56, 122-123.	0.3	18
454	Haematological abnormalities in mitochondrial disorders. <i>Singapore Medical Journal</i> , 2015, 56, 412-419.	0.3	15
455	Dilative Arteriopathy and Leucoencephalopathy as Manifestations of a Neurometabolic Disease. <i>The Open Neurology Journal</i> , 2015, 9, 28-31.	0.4	7
456	Cardiac manifestations of primary mitochondrial disorders. <i>International Journal of Cardiology</i> , 2014, 177, 754-763.	0.8	95
457	CNS disease triggering Takotsubo stress cardiomyopathy. <i>International Journal of Cardiology</i> , 2014, 177, 322-329.	0.8	125
458	Treatment of dystrophin cardiomyopathies. <i>Nature Reviews Cardiology</i> , 2014, 11, 168-179.	6.1	57
459	CNS-disease affecting the heart: Brain-heart disorders. <i>Journal of the Neurological Sciences</i> , 2014, 345, 8-14.	0.3	55
460	Anterocollis and anterocaput. <i>Clinical Neurology and Neurosurgery</i> , 2014, 127, 44-53.	0.6	8
461	Asymptomatic neurocysticercosis over 15 years between transitory ischemic attack as initial and epilepsy as second manifestation. <i>Acta Neurologica Belgica</i> , 2013, 113, 203-204.	0.5	1
462	Influence of mitochondrion-toxic agents on the cardiovascular system. <i>Regulatory Toxicology and Pharmacology</i> , 2013, 67, 434-445.	1.3	66
463	Presentation of adult mitochondrial epilepsy. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2013, 22, 119-123.	0.9	20
464	Noncompaction predisposing for recurrent Takotsubo syndrome in myotonic dystrophy 1. <i>International Journal of Cardiology</i> , 2013, 168, e38-e39.	0.8	13
465	Pathomorphologic findings in left ventricular hypertrabeculation/noncompaction of adults in relation to neuromuscular disorders. <i>International Journal of Cardiology</i> , 2013, 169, 249-253.	0.8	31
466	CRYPTOGENIC VITAMIN-D-DEFICIENCY MANIFESTING AS GENERALISED FASCICULATIONS AND CARPOPEDAL SPASMS. <i>Acta Clinica Belgica</i> , 2013, 68, 451-452.	0.5	1
467	Mitochondrial epilepsy in pediatric and adult patients. <i>Acta Neurologica Scandinavica</i> , 2013, 128, 141-152.	1.0	16
468	Increased prevalence of malignancy in adult mitochondrial disorders. <i>Journal of Medicine and Life</i> , 2013, 6, 477-81.	0.4	9

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469	Mitochondrial toxicity of antiepileptic drugs and their tolerability in mitochondrial disorders. Expert Opinion on Drug Metabolism and Toxicology, 2012, 8, 71-79.	1.5	93
470	Epilepsy in mitochondrial disorders. Seizure: the Journal of the British Epilepsy Association, 2012, 21, 316-321.	0.9	73
471	Primary mitochondrial arteriopathy. Nutrition, Metabolism and Cardiovascular Diseases, 2012, 22, 393-399.	1.1	17
472	Acquired, familial noncompaction and eccentric hypertrophic cardiomyopathy associated with metabolic myopathy and epilepsy. International Journal of Cardiology, 2012, 160, 73-75.	0.8	11
473	Inherited Mitochondrial Disorders. Advances in Experimental Medicine and Biology, 2012, 942, 187-213.	0.8	34
474	Do pets reduce the likelihood of sudden unexplained death in epilepsy?. Seizure: the Journal of the British Epilepsy Association, 2012, 21, 649-651.	0.9	13
475	Stroke and Stroke-like Episodes in Muscle Disease. The Open Neurology Journal, 2012, 6, 26-36.	0.4	36
476	Mimicry between mitochondrial disorder and multiple sclerosis. Metabolic Brain Disease, 2012, 27, 217-220.	1.4	5
477	Inherited mitochondrial neuropathies. Journal of the Neurological Sciences, 2011, 304, 9-16.	0.3	52
478	Left ventricular hypertrabeculation/noncompaction in hereditary inclusion body myopathy. International Journal of Cardiology, 2011, 150, e67-e69.	0.8	13
479	Treatment of central nervous system manifestations in mitochondrial disorders. European Journal of Neurology, 2011, 18, 28-38.	1.7	23
480	Frequency of Stroke and Embolism in Left Ventricular Hypertrabeculation/Noncompaction. American Journal of Cardiology, 2011, 108, 1021-1023.	0.7	129
481	Parkinson's syndrome and Parkinson's disease in mitochondrial disorders. Movement Disorders, 2011, 26, 784-791.	2.2	23
482	Transition m.3308T>C in the ND1 Gene Is Associated with Left Ventricular Hypertrabeculation/Noncompaction. Cardiology, 2011, 118, 153-158.	0.6	17
483	Orthopaedic abnormalities in primary myopathies. Acta Orthopaedica Belgica, 2011, 77, 563-82.	0.1	16
484	Left ventricular non-compaction and its cardiac and neurologic implications. Heart Failure Reviews, 2010, 15, 589-603.	1.7	38
485	Perspectives of Kennedy's disease. Journal of the Neurological Sciences, 2010, 298, 1-10.	0.3	121
486	Acquired noncompaction associated with coronary heart disease and myopathy. Heart and Lung: Journal of Acute and Critical Care, 2010, 39, 240-241.	0.8	16

#	ARTICLE	IF	CITATIONS
487	Successful low-dose azathioprine for myasthenia gravis despite hepatopathy from primary sclerosing cholangitis: a case report. <i>Journal of Medical Case Reports</i> , 2010, 4, 356.	0.4	3
488	Mitochondrial disorder mimicking ocular myasthenia. <i>Acta Neurologica Belgica</i> , 2010, 110, 110-2.	0.5	9
489	Is mitochondrial disease the common cause of histiocytoid cardiomyopathy and non-compaction?. <i>International Journal of Legal Medicine</i> , 2009, 123, 507-508.	1.2	7
490	Cardiogenetics, Neurogenetics, and Pathogenetics of Left Ventricular Hypertrabeculation/Noncompaction. <i>Pediatric Cardiology</i> , 2009, 30, 659-681.	0.6	219
491	Bulbar and spinal muscular atrophy (Kennedy's disease): a review. <i>European Journal of Neurology</i> , 2009, 16, 556-561.	1.7	67
492	Management of mitochondrial stroke-like episodes. <i>European Journal of Neurology</i> , 2009, 16, 1178-1184.	1.7	36
493	Mitochondrial Ataxias. <i>Canadian Journal of Neurological Sciences</i> , 2009, 36, 543-553.	0.3	34
494	Central Nervous System Imaging in Mitochondrial Disorders. <i>Canadian Journal of Neurological Sciences</i> , 2009, 36, 143-153.	0.3	29
495	Systemic and non-systemic vasculitis affecting the peripheral nerves. <i>Acta Neurologica Belgica</i> , 2009, 109, 100-13.	0.5	14
496	Management of peripheral facial nerve palsy. <i>European Archives of Oto-Rhino-Laryngology</i> , 2008, 265, 743-752.	0.8	224
497	Cognitive decline as a manifestation of mitochondrial disorders (mitochondrial dementia). <i>Journal of the Neurological Sciences</i> , 2008, 272, 20-33.	0.3	51
498	Atrial fibrillation/flutter in myopathies. <i>International Journal of Cardiology</i> , 2008, 128, 304-310.	0.8	35
499	Leigh and Leigh-Like Syndrome in Children and Adults. <i>Pediatric Neurology</i> , 2008, 39, 223-235.	1.0	330
500	Don't miss extra-cardiac manifestations of familial left ventricular hypertrabeculation/non-compact. <i>European Journal of Heart Failure</i> , 2007, 9, 101-102.	2.9	0
501	Hematological Manifestations of Primary Mitochondrial Disorders. <i>Acta Haematologica</i> , 2007, 118, 88-98.	0.7	71
502	Is atherosclerosis a mitochondrial disorder?. <i>Vasa - European Journal of Vascular Medicine</i> , 2007, 36, 229-240.	0.6	25
503	CONSEQUENCES OF MISDIAGNOSING MITOCHONDRIAL DISORDER. <i>International Journal of Neuroscience</i> , 2006, 116, 907-914.	0.8	3
504	Is myocardial damage truly absent in inclusion body myositis with elevated troponin T level?. <i>Human Pathology</i> , 2006, 37, 1367-1368.	1.1	2

#	ARTICLE	IF	CITATIONS
505	Mitochondrial Disorder Aggravated by Propranolol. <i>Southern Medical Journal</i> , 2006, 99, 768-771.	0.3	11
506	Madarosis from mitochondriopathy. <i>Acta Ophthalmologica</i> , 2005, 83, 628-630.	0.4	6
507	Left Ventricular Hypertrabeculation/Noncompaction and Stroke or Embolism. <i>Cardiology</i> , 2005, 103, 68-72.	0.6	87
508	Mitochondrial neuropathy. <i>Clinical Neurology and Neurosurgery</i> , 2005, 107, 181-186.	0.6	36
509	Left ventricular hypertrabeculation/noncompaction as a cardiac manifestation of Duchenne muscular dystrophy under non-invasive positive-pressure ventilation. <i>Acta Cardiologica</i> , 2005, 60, 445-448.	0.3	26
510	Stress lactate in mitochondrial myopathy under constant, unadjusted workload. <i>European Journal of Neurology</i> , 2004, 11, 811-816.	1.7	27
511	Lone noncompaction in Leber's hereditary optic neuropathy. <i>Acta Cardiologica</i> , 2004, 59, 187-190.	0.3	12
512	Mitochondriopathy as a differential diagnosis of amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases</i> , 2002, 3, 219-224.	1.4	17
513	Left ventricular hypertrabeculation/noncompaction and association with additional cardiac abnormalities and neuromuscular disorders. <i>American Journal of Cardiology</i> , 2002, 90, 899-902.	0.7	523
514	Phenotype variability in 130 adult patients with respiratory chain disorders. <i>Journal of Inherited Metabolic Disease</i> , 2001, 24, 560-576.	1.7	53
515	Wolff-Parkinson-White syndrome and isolated left ventricular abnormal trabeculation as a manifestation of Leber's hereditary optic neuropathy. <i>Canadian Journal of Cardiology</i> , 2001, 17, 464-6.	0.8	21
516	Complex Mitochondriopathy Associated with 4 mtDNA Transitions. <i>European Neurology</i> , 2000, 44, 37-41.	0.6	46
517	The Apparent Beneficial Effect of L-arginine for Stroke-like Lesions can Be Accidental. <i>Neurohospitalist, The</i> , 0, , 194187442210897.	0.3	0
518	Discrimination between Benign and Malignant Post-SARS-CoV-2 Vaccination Lymphadenopathy is Feasible. <i>Korean Journal of Radiology</i> , 0, 23, .	1.5	1
519	Take children with progressive quadruparesis after SARS-CoV-2 infection seriously. <i>Child's Nervous System</i> , 0, , .	0.6	0
520	Letter to the Editor Regarding "Colchicine Against SARS-CoV-2 Infection: What is the Evidence?" <i>Rheumatology and Therapy</i> , 0, , .	1.1	0
521	Rule Out Alternative Causes Before Attributing Acute, Haemorrhagic, Leukoencephalitis (AHLE) to COVID-19. <i>Neurohospitalist, The</i> , 0, , 194187442211103.	0.3	0
522	Commentary: Point Prevalence and Associated Factors of Hip Displacement in Pediatric Patients With Mitochondrial Disease. <i>Frontiers in Pediatrics</i> , 0, 10, .	0.9	0

#	ARTICLE	IF	CITATIONS
523	Neurological Side Effects of SARS-CoV-2 Vaccinations are Not Uncommon but Frequently Ignored [Letter]. Neuropsychiatric Disease and Treatment, 0, Volume 18, 1273-1275.	1.0	2
524	Before blaming a COVID vaccine for cytotoxic lesions of the corpus callosum all other differentials must be ruled out. Neuroradiology, 0, , .	1.1	1
525	Letter to the Editor regarding "COVID-19-Related Burden and Risk Perception in Individuals with Chronic Inflammatory Demyelinating Polyneuropathy and Multifocal Motor Neuropathy: A Cross-Sectional Study" Neurology and Therapy, 0, , .	1.4	0